

K112389.

**510(k) Summary**

JUL 20 2012

**Submitter information**

<b>Company name</b>	Materialise N.V.
<b>Establishment registration number</b>	3003998208
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<b>Contact name</b>	Alexandra Razzhivina
<b>Contact title</b>	Regulatory officer
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**Submission date**

The date of the Traditional 510(k) submission is 16<sup>th</sup> of August, 2011.

**Submission information**

<b>Trade Name</b>	SurgiCase Orthopaedics, SurgiCase Connect, SurgiCase Guides
<b>Common Name</b>	Image processing system and software for simulating/evaluating implant placement and surgical treatment options
<b>Classification Name</b>	Radiological image processing system
<b>Product code</b>	PBF

**Predicate devices**

<b>Trade or proprietary or model name</b>	SurgiCase
<b>510(k) number</b>	K073449
<b>Decision date</b>	2008/04/16
<b>Product code</b>	LLZ
<b>Manufacturer</b>	Materialise N.V.

<b>Trade or proprietary or model name</b>	SurgiCase Guide
<b>510(k) number</b>	K103136
<b>Decision date</b>	2011/04/18
<b>Product code</b>	JEY, MQN
<b>Manufacturer</b>	Materialise N.V.

## ***Device Information***

### **Description and functioning of the device**

The SurgiCase Orthopaedics system is composed of two components: SurgiCase Connect (software) and SurgiCase Guides (hardware).

The SurgiCase Orthopaedics system is intended to be used as a surgical instrument to transfer a pre-surgical plan to the lower and upper extremity during orthopaedic surgical procedures.

SurgiCase Connect is a medical device for Materialise and a surgeon for pre-surgical simulation and evaluation of surgical treatment options. This includes transferring, visualizing, measuring, annotating and editing medical data.

The SurgiCase Guides are patient specific templates that are based on a pre-surgical software plan and are designed to fit a specific patient. All guides are individually designed and manufactured for each patient using a design and manufacturing process with strict procedures and work instructions to guarantee guides that consistently perform in a safe and effective way. In surgery these guides are used to assist a surgeon in guiding the marking of bone and/or guiding surgical instruments according to the pre-surgical plan.

### ***Intended use***

The SurgiCase Orthopaedics system is intended to be used as a surgical instrument to assist in pre-operative planning and/or in guiding the marking of bone and/or guide surgical instruments in non-acute, non-joint replacing osteotomies for upper extremity orthopedic surgical procedures.

The system is to be used for adult patients.

SurgiCase Guides are intended for single use only.

### ***Summary of technological characteristics***

Device comparison showed that the proposed device is substantially equivalent in intended use, materials and performance characteristics to the predicate devices.

### ***Performance data***

#### **Non-clinical tests**

The SurgiCase Connect software has been validated for its intended use to determine substantial equivalence to the predicate devices.

SurgiCase Guides were validated through non-clinical studies using bone models and cadaver specimens. Biocompatibility test, sterility test, sterilization dimensional stability test, debris test and packaging and shipment test and cleaning validation test were performed to assess the safety and effectiveness of the SurgiCase Guides. Testing verified that the accuracy and performance of the device is adequate to perform as intended.



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration  
10903 New Hampshire Avenue  
Document Control Room –WO66-G609  
Silver Spring, MD 20993-0002

Materialise N.V.  
% Ms. Alexandra Razzhivina  
Technologielaan 15  
Leuven, Belgium 3001

JUL 30 2012

Re: K112389

Trade/Device Name: Surgicase Connect, Surgicase Guide

Regulation Number: 21 CFR 888.3030

Regulation Name: Single/multiple component metallic bone fixation appliances and accessories

Regulatory Class: II

Product Code: PBF

Dated: July 06, 2012

Received: July 09, 2012

Dear Ms. Razzhivina:

This letter corrects our substantially equivalent letter of July 20, 2012 for the product code and regulation number for your medical device. We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); medical

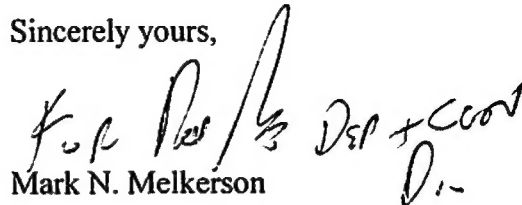
Page 2 – Ms. Alexandra Razzhivina

device reporting (reporting of medical device-related adverse events) (21 CFR 803); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please go to <http://www.fda.gov/AboutFDA/CentersOffices/CDRH/CDRHOffices/ucm115809.htm> for the Center for Devices and Radiological Health's (CDRH's) Office of Compliance. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to <http://www.fda.gov/MedicalDevices/Safety/ReportaProblem/default.htm> for the CDRH's Office of Surveillance and Biometrics/Division of Postmarket Surveillance.

You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address <http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm>.

Sincerely yours,

A handwritten signature in black ink, appearing to read "Mark N. Melkerson" with a stylized flourish at the end.

Mark N. Melkerson

Director

Division of General, Restorative  
and Neurological Devices  
Office of Device Evaluation  
Center for Devices and  
Radiological Health

## Indications for Use

510(k) Number (if known): K112389

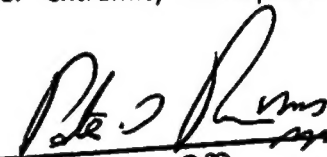
Device Name: The SurgiCase Orthopaedics system

### Indications for Use:

The SurgiCase Orthopaedics system is intended to be used as a surgical instrument to assist in pre-operative planning and/or in guiding the marking of bone and/or guide surgical instruments in non-acute, non-joint replacing osteotomies for upper extremity orthopedic surgical procedures.

The system is to be used for adult patients.

SurgiCase Guides are intended for single use only.

  
(Division Sign-Off)  
Division of Surgical, Orthopedic,  
and Restorative Devices

510(k) Number K112389

Prescription Use   X    
(Part 21 CFR 801 Subpart D)

AND/OR

Over-The-Counter Use         
(21 CFR 801 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF  
NEEDED)

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Concurrence of CDRH, Office of Device Evaluation (ODE)